QUARTERLY GROUNDWATER MONITORING REPORT

FOR

FORMER ANGELES CHEMICAL COMPANY FACILITY 8915 SORENSEN AVENUE SANTA FE SPRINGS, CALIFORNIA

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1.0) INTRODUCTION

Blakely Environmental Investigations, Inc. (BEII) was contracted by Greve Financial Services ((310) 753-5770) to perform quarterly groundwater monitoring at the former Angeles Chemical Company, Inc. facility located at 8915 Sorensen Avenue, Santa Fe Springs, California (See Figure 1, Site Location Map). The quarterly groundwater monitoring was requested by the Department of Toxics Substance Control (DTSC) correspondence dated September 18, 2001.

2.0) SITE LOCATION AND HISTORY

The site is approximately 1.8 acres in size and completely fenced. The site was bound to Sorensen Avenue on the east, Liquid Air Corporation to the northwest, Plastall Metals Corporation to the north, and a Southern Pacific Railroad easement and Mckesson Chemical Company to the south.

The property was owned by Southern Pacific Transportation Company and was not developed until 1976.

The Angeles Chemical Company has operated as a chemical repackaging facility since 1976. A total of thirty-four (34) underground storage tanks (USTs) existed beneath the site. Two (2) USTs, one gasoline and one diesel, and sixteen (16) chemical USTs were excavated and removed under the oversight of the Santa Fe Springs Fire Department. All 16 remaining chemical USTs were decommissioned in place and slurry filled.

In January 1990, SCS Engineers, Inc. (SCS) conducted a site investigation. SCS advanced eight borings from 5' below grade (bg) to 50' bg. Soil samples collected and analyzed identified benzene, 1,1-Dichloroethane (1,1-DCA), 1,1-Dichloroethene (1,1-DCE), MEK, methyl isobutyl ketone (MIBK), toluene, 1,1,1-TCA, PCE, and xylenes at detectable concentrations.

In June 1990, SCS performed an additional site investigation at the site by advancing six additional borings advanced from 20.5' bg to 60' bg. A monitoring well (MW-1) was also installed. Soil sample analysis identified detectable concentrations of the above mentioned VOCs in addition to acctone and methylene chloride. Dissolved benzene, 1,1-DCA, 1,1-DCE, PCE, TCE, and trans-1,2-dichloroethene were detected in MW-1 above maximum contaminant levels.

Between 1993 and 1994, SCS performed further testing at the site. Soil samples were collected from nine borings. Five borings were converted to groundwater monitoring wells MW-2, MW-3, MW-4, MW-6, and MW-7 (See Figure 2, Well Location Map). The predominant compounds detected in soil were acetone, MEK, MIBK, PCE, toluene, 1,1,1-TCA, TCE, and xylenes. Groundwater sample collection performed in

February 1994 by SCS identified the following using EPA method 624 (laboratory results included in Remedial Investigation Report dated August 1994 by SCS):

			General Color of American Color			en e
1,1-DCA	649	1,130	85	1,410	2,260	2,130
L2-DCA	<100			<u>∻~⊬⊀100°%∨</u>	1,140	31- × 151
1,1-DCE	2,210	2,460	2,800	806 33311-1 30 7-33	1,240 1,910	
Methylene Chloride	1,220	2,980	6,530	4,760	21,400	<30
"是这种的人,是是不是是是是一个人,	**************************************	2110	** 5370 **/	3,320 12,700	12 500	134 398
Toluene	560	7,390	579	12,700	13,500	290 at
TCE	7,160	3,040	1,730	14,300	1,320	45
		7.70	**************************************	THE THE	4,110	1.06
Units	μ e/L	μg/L	μ <u>e</u> /L	μg/L	<u>μα/L</u>	<u>μ</u> με/Σ

In 1996, SCS performed separate soil vapor extraction pilot testing beneath the site at approximately 10' bg and 22' bg. Laboratory analysis identified maximum soil vapor gas concentrations as 1,1,1-TCA (30,300 ppmV) with detectable concentrations of 1,1-DCE, TCE, methylene chloride, toluene, PCE and xylenes. The maximum radius of influence from the various extraction units used were measured as 35 feet at 10' bg and 80 feet at 22' bg.

In November 1997, SCS performed a soil vapor survey at the site. Soil vapor samples were collected at twenty-three locations at 5' bg. In addition, soil vapor samples were collected at 15' bg in five of the twelve sampling points. The soil vapor survey identified maximum VOC contaminants near the railroad tracks on site, the location where a rail tanker reportedly had an accidental release.

In July 2000, BEII contracted BLC Surveying, Inc. to perform a site survey. Well locations were recorded using the California Plane coordinate systems. A copy of the survey is on file with the DTSC.

In September 2000, Blaine Tech Services, Inc. gauged the six on-site monitoring wells (MW-1, MW-2, MW-3, MW-4, MW-6, and MW-7) under the supervision of BEII. Free product (FP) was identified in monitoring well MW-4 at 0.21-feet in thickness. Approximately 0.5 liters of FP were removed from the well and placed in a scaled 55-gallon drum.

BEII performed a soil vapor gas survey at the site from November 27 to December 1, 2000. A total of 36 soil vapor sample points, labeled SV1 through SV36, were selected by BEII and approved by the DTSC for analysis. Two discrete soil vapor samples were collected from each soil vapor sample point, one at 8' bg and one at 20' bg. SV1 was an exception since the first soil vapor sample was collected at 10' bg instead of

8' bg (See Figure 3 for BEII Soil Vapor Sample Locations). Based on the soil vapor sample results, BEII identified relatively low level concentrations of VOCs in the silty clay soils at 8' bg. However, the concentrations of VOCs are significantly higher in the sandy soils at 20' bg in OU-1. Results were submitted to the DTSC by BEII in a Report of Findings dated January 10, 2001 with laboratory reports (BEII Report of Findings dated January 10, 2001).

On November 30, 2000, Blaine Tech Services, Inc. (Blaine) was contracted to perform groundwater sampling at the site. Groundwater monitoring wells MW-4 and MW-6 identified were not sampled due to insufficient water and presence of free product. These wells were installed to monitor a perched groundwater body to the north. Free product was identified in MW-1 during sample collection, upon completion of well purging. The potentiometric groundwater level was above the well screen. Groundwater purging lowered the potentiometric level below the screened interval, allowing free product to enter. Groundwater sample analysis identified thirteen constituents of concern (COCs) in the dissolved phase as VOCs only. Laboratory analysis of metals and SVOCs identified concentrations below allowable levels for those constituents. Results were submitted by BEII to the DTSC in a Report of Findings dated January 10, 2001 with laboratory reports.

3.0) REGIONAL GEOLOGY/HYDROGEOLOGY

The site is located near the northern boundary of the Santa Fe Springs Plain within the Los Angeles Coastal Plain at an elevation of approximately 150 feet above mean sea level. Surficial sediments consist of fluvial deposits composed of inter-bedded gravel, sand, silt, and clay. Available data from California Water Resources Bulletin No. 104 (June 1961) indicate that the surficial sediments may be Holocene and/or part of the upper Pleistocene Lakewood Formation, which ranges from 40 to 50 feet thick beneath the site. The Lakewood Formation has lateral lithologic changes with discontinuous permeable zones that vary in particle size. Stratified deposits of sand, silty sand, silt, and fine gravel comprising the upper portion of the lower Pleistocene San Pedro Formation underlies the Lakewood Formation.

The site lies within the Central Basin Pressure area, a division of the Central Ground Water Basin, which extends over most of the Coastal Plain. The Gasper aquifer, a part of the basal coarse unit of Holocene deposits, is found within old channels of the San Gabriel and other rivers. The Gasper aquifer may be 40-feet in thickness, with its base at a depth of about 80 to 100-feet bg. The underlying Gage aquifer is found within the upper Pleistocene Lakewood Formation. The Hollydale aquifer is the uppermost regional aquifer in the San Pedro Formation. Bulletin 104 indicates that this aquifer averages approximately 30-feet in thickness in this area, with its top at a depth of about 70 feet bg. The major water producing aquifers in the region are the Lynwood aquifer located approximately 200-feet bg, the Silverado aquifer located at approximately 275-feet bg, and the Sunnyside aquifer located at approximately 600-feet bg.

4.0) SITE GEOLOGY/HYDROGEOLOGY

SCS identified silty clays with some minor amounts of silt and sand in the shallow subsurface from surface grade to approximately 15' bg. Below the silty clay, poorly sorted coarse-grained sand and gravel from 15' bg to 26' bg. A less permeable silty clay layer was identified by SCS between 35' and 50' bg, which contains stringers of fine sand and silt that is part of the Gaspur/Hollydale aquifer.

Two aquifers were identified by SCS during subsurface investigations performed at the site. A perched aquifer was encountered at approximately 23' bg and the Gaspur/Hollydale aquifer was encountered at 20' to 35' bg by SCS. The groundwater gradient flows to the southwest as identified by SCS. In October 2001, the groundwater was identified between 26.35' bg to 39.19' bg beneath the site with a flow to the south/southwest (See Figure 4, Groundwater Gradient Map). Only monitoring wells installed in the Gasper/Hollydale aquifer (MW-1, MW-2, MW-3, and MW-7) were used to calculate the groundwater gradient.

5.0) GROUNDWATER MONITORING PROTOCOL

The purpose of the proposed groundwater monitoring was to provide data regarding the piezometric surface, water quality, and the presence of free product (FP), if any on a quarterly basis to the DTSC. Groundwater monitoring consisted of such activities as water level measurement, well sounding for detection of FP, collection of groundwater samples, field analysis, laboratory analysis, and reporting. The proposed work was performed as follows:

The depth to groundwater was measured in each well using a decontaminated water level indicator capable of measuring to with 1/100th of a foot. Prior to and following collection of measurements from each well, the portions of the water level indicator entering groundwater were decontaminated using a 3-stage decontamination consisting of a potable wash with water containing Liquinox soap followed by a double purified water rinse. Wells were monitored in the order of least contaminated to the most contaminated based on past analysis. For the Angeles Chemical Co. wells, the following order of wells was followed: MW-7, MW-3, MW-2, MW-1, MW-4, and MW-6.

The well box and easing were opened carefully to minimize the amount of debris or dirt falling into the open easing. Once the well cap and J-Plug were removed, the water level indicator was lowered into the well until a consistent tone is registered. Several soundings were repeated to verify the measured depth to groundwater. The depth of groundwater was measured from a reference point marked on the lip of each well casing. A licensed surveyor has surveyed the elevation of each reference point. The result was recorded on the field sampling log for each well. Other relevant information such as physical condition of the well, presence of hydrocarbon odors, etc. was also recorded as appropriate on the field sampling log.

The weil sounder used for this project will also be equipped to measure FP layers thicker than 0.1 inches. FP was indicated as light non-aqueous phase liquid (LNAPL) or dense non-aqueous phase liquid (DNAPL).

Groundwater sampling was conducted immediately following the collection of groundwater depth measurements from all monitoring wells. Free product present within any well was sampled and characterized upon completion of groundwater sampling to minimize cross contamination.

Groundwater samples were analyzed for the following constituents:

 Volatile organic compounds (VOCs) using EPA Method 8260B to include all Tentatively Identified Compounds (TICs).

 Title 22 (CAM 17 heavy metals) metals using EPA Method 6010 and 7471 for mercury.

5.1) Well Purging and Measurement of Field Parameters

Wells were purged in the following order MW-7, MW-3, MW-2, and MW-1 to minimize the potential for cross contamination. The wells were purged by Blaine Tech Services, Inc (Blaine) and sampled by BEII on October 30, 2001 in the presence of Mr. Ryan Kinsella and Mr. Sanford Britt of the DTSC. The purge protocol was presented in the Field Sampling Plan as Appendix A in the Groundwater Monitoring Work Plan dated October 23, 2001.

Prior to purging, casing volumes was calculated based on total well depth, standing water level, and casing diameter. One casing volume was calculated as:

$$V = \pi (d/2)^2 h \times 7.48$$

where:

V is the volume of one well casing of water (in gallons, $1 \text{ ft}^3 = 7.48$ gallon);

d is the inner diameter of the well casing (in feet); and

h is the total depth of water in the well - the depth to water level (in feet).

A minimum of three casing volumes of water was purged unless wells were de-watered, as in the case of monitoring wells MW-3 and MW-2. Only 17-gallons were purged from MW-3 and MW-2 because the wells observed a slow recharge. Water was collected into a measured bucket to record the purge volume. All purged groundwater was containerized in 55-gallon hazardous waste drum for disposal at a later date.

After each well casing volume was purged; water temperature, pH,

specific conductance (EC), and turbidity were measured using field test meters and the measurements were recorded on Well Monitoring Data Sheets (See Appendix A). Samples were collected after these parameters have stabilized; indicating that representative formation water has entered the well. The temperature, pH, and specific conductance should not vary by more than 10 percent from reading to reading. Turbidity should be less then 5 NTUs, however, the purging process stirred up silty material in each well which made the turbidity measurements of 5 NTUs unattainable. Notations of water quality including color, clarity, odors, sediment, etc. were also noted in the data sheets.

Groundwater samples were collected after water levels recharged to 80 percent of the static water column, with the exception of MW-3. The groundwater level in MW-3 recharged to only 60 percent when the sample was collected 3-hours after purging.

All field meters were calibrated according to manufacturers' guidelines and specifications before and after each day of field use. Field meter probes were decontaminated before and after use at each well. The pH, conductivity, and temperature were measured with a Myron-L Ultra Meter and turbidity was measured with a HF Scientific DRT-15C meter. The calibration standards used for pH were 4 and 7 with expiration dates of December 2001. Conductivity was calibrated to a 3900 µs standard with an expiration date of January 2002. A 0.02 NTU standard was used to calibrate the turbidity with an expiration date of January 2002.

5.2) Weil Sampling

Groundwater and FP samples were collected by lowering a disposable bailer into each well. Groundwater and FP were transferred from the bailer directly into the appropriate sample containers with preservative, if required, chilled, and processed for shipment to the laboratory. When transferring samples, care was taken not to touch the bailer-emptying device to the sample containers. Water and FP samples were transported to Southland Technical Services, Inc., a certified laboratory by the California Department of Health Services (Cert. #1986) to perform the requested analysis.

Groundwater samples were collected from monitoring wells MW-7, MW-3, MW-2, and MW-1 only. Monitoring wells MW-4 and MW-6 identified FP as LNAPL at a thickness of 0.3' and 0.5', respectively. FP was collected in a 40 ml VOA for laboratory analysis from only MW-6. Monitoring well MW-4 contained insufficient liquids to obtain a FP sample. FP collected from MW-6 was analyzed for Volatile organic compounds (VOCs) using EPA Method 8260B to include all Tentatively Identified Compounds (TICs). The FP sample was collected after all groundwater samples were collected to minimize cross contamination.

Vials for VOC analysis were filled first to minimize aeration of groundwater collected in the bailer. The laboratory provided vials containing sufficient HCl preservative to lower the pH to less than 2. The vials were filled directly from the bottom-emptying device. The vial was capped with a cap containing a Teflon septum. The vial was inverted and tapped to check for bubbles to insure zero headspace.

Groundwater sample collection for dissolved metals followed. The laboratory provided 500-mL poly containers containing sufficient HNO₃ preservative. The containers were filled directly from the bottom-emptying device.

New nitrile gloves were worn during by sampling personnel for each well to prevent cross contamination of the samples. A solvent free label was affixed to each sample container/vial denoting the well identification, date and time of sampling, and an identifying code to distinguish each individual bottle.

5.3) Sample Handling

VOA vials and 500-mL poly containers, including laboratory trip blanks and rinse blanks, were placed inside of one new Ziplock bag per well and stored in a cooler chilled to approximately 4°C with bagged ice. Water and FP samples were logged on the chain-of-custody forms immediately following sampling of each well to insure proper tracking through analysis to the laboratory.

5.4) Waste Management

FP, purged groundwater, and decontamination water were stored in sealed 55-gallon drums for a period not exceeding 90 days. Stored wastes will be profiled for hazardous constituents and characterized as Non-Hazardous, California Hazardous, or RCRA Hazardous, as appropriate. Any transportation of waste will be under appropriate manifest.

6.0) FP AND GROUNDWATER SAMPLE RESULTS

Laboratory analysis of FP collected from perched water monitoring well MW-6 identified 1,1,1 TCA at 28,100 mg/L (2.8%), 1,2,4-Trimethylbenzene at 22,100 mg/L (2.2%), Xylenes at 10,370 mg/L (1.0%), Toluene at 9,010 mg/L (0.9%), 1,3,5-Trimethylbenzene at 5,400 mg/L (0.5%), and Ethylbenzene at 4,320 mg/L (0.4%). Other VOCs were detected in the free product at concentrations less than 2,000 mg/L or 0.2% and are listed in Table 1. Laboratory groundwater and FP analysis reports are included as Appendix B.

Groundwater sample results from the Gasper/Hollydale aquifer identified relatively low VOC concentrations to the north end of the site and relatively high VOC concentrations to the south along the McKesson property (See Table 1).

Dissolved BTEX concentrations were identified as 1,438 µg/L in MW-7 located along the northern side of the property. On the southern side of the property, dissolved BTEX concentrations ranged from 10,530 µg/L to 328 µg/L in monitoring wells MW-1 through MW-3 (See Figure 5, Estimated Dissolved BTEX Plume). Most of the dissolved BTEX concentrations from all Gasper/Hollydale monitoring wells consisted of toluene and xylene as 84-89% of the total BTEX, with the exception of MW-2 which identified mostly benzene and ethylbenzene as 92% of the total BTEX. However, monitoring well MW-2 contained the least amount of total dissolved BTEX at 328 µg/L.

Dissolved PCE was identified in monitoring wells MW-3 and MW-7 exclusively. Monitoring well MW-3 identified dissolved PCE at 130 µg/L and well MW-7 identified dissolved PCE at 100 µg/L (See Figure 6 for Estimated Dissolved PCE Plume). Dissolved TCE was identified exclusively in MW-3 at 100 µg/L (See Figure 7 for Estimated Dissolved TCE Plume). Again, dissolved VOC concentrations were detected at higher levels along the southern side of the property.

Concentrations of chlorinated VOC daughter products also show a trend of relatively high dissolved concentrations along the southern side of the property. 1,1 DCA is a daughter product from reductive dehalogenation of 1,1,1-TCA and from carbon-carbon double bond reduction of 1,1 DCE, another daughter product. Dissolved 1,1 DCA was identified at a concentration of 2,670 µg/L in MW-7, which is located along the northern side of the property. Concentrations of dissolved 1,1 DCA along the southern side of the property were identified as 8,190 µg/L in MW-1, 1,500 µg/L in MW-2, and 1,030 µg/L in MW-3 (See Figure 8 for Estimated Dissolved 1,1 DCA Plume).

Dissolved 1,1 DCE, a daughter product of the dehydrohalogenation of 1,1,1 TCA and reductive dehalogenation of TCE, was also identified at a concentration of 355 µg/L in MW-7. Along the southern side of the property, dissolved 1,1 DCE was identified in wells MW-1 through MW-3 at concentrations of 4,090 µg/L to 1,120 µg/L (See Figure 9 for Estimated Dissolved 1,1 DCE Phune). Concentrations of dissolved 1,1 DCE were 3 to 11 times greater along the southern side of the property compared to the northern side.

Cis-1,2 DCE is also a daughter product of the dehydrohalogenation of 1,1,1 TCA and reductive dehalogenation of TCE. Concentrations of dissolved cis-1,2 DCE were identified along the north side of the property at 194 µg/L. The southern side of the property identified dissolved cis-1,2 DCE from 10,300 µg/L to 7,000 µg/L in monitoring wells MW-1 through MW-3 (See Figure 10 for Estimated Dissolved cis-1,2 DCE Plume). Dissolved cis-1,2 DCE concentrations were 36 to 53 times greater along the southern side of the property compared to the northern side.

Vinyl chloride (VC) is a by product from the dehydrohalogenation and reductive dehalogenation of the chlorinated VOC daughter products mentioned above. Dissolved VC concentrations were identified along the southern side of the property in monitoring wells MW-1 and MW-2 at 1,350 μ g/L and 75 μ g/L, respectively (See Figure 11 for Estimated Dissolved VC Plume). Concentrations of dissolved VC were identified along the northern side of the property in MW-7 at 188 μ g/L. Again, dissolved VOC concentrations were detected at higher levels along the southern side of the property.

Groundwater samples were filtered by the laboratory and analyzed for dissolved metals (See Table 2 for Dissolved Metal Results). Arsenic was the only metal that was identified above the maximum contaminant level (MCL) of 50 μ g/L. Arsenic was identified as 61 μ g/L in MW-2 and 71 μ g/L in MW-7.

7.0) CONCLUSIONS

Based on the significant increase in concentrations of dissolved VOCs confined to the south side of the site, BEII concludes that the probable source of the majority of groundwater VOC impact is located south of the former Angeles Chemical Co. property. It is concluded that the dissolved groundwater plume has migrated onto the Angeles site in the presently dry shallow permeable soil.

This permeable soil was identified as saturated with water and DNAPLs during from 1975 until 1986 when McKesson dumped DNAPL /wastewater mix into the unlined ditch adjoining the south side of the former Angeles Chemical property. It was identified by the USEPA in 1986 that 1,500 gallons per day of wastewater with approximately 1.5% DNAPL were continuously discharged in these areas, resulting in saturation of the shallow permeable zone from 1975 until 1986.

It is further concluded that this large source release into groundwater created a hydraulic pressure head driving the DNAPL/water from surface through the permeable soil to the low point at MW-6, approximately 30' bg, which also provided communication with the Gasper/Hollydale Aquifer now being tested (See Figure 12).

It is further concluded that the DNAPL saturated soil continues to act as a source beneath the unlined ditch and bermed discharge area of McKesson and has diffused as a concentration gradient from high levels at the source to low levels away from the source with residual DNAPL, LNAPL and water pooling at the lowest point of the permeable soil zone identified by monitoring well MW-6. Saturated flow of solvents identified in the shallow permeable soil of the McKesson site from 15' bg to 30' bg and sloping from the unlined ditch onto the former Angeles Chemical property has impacted the lowest measured point of this continuous permeable layer, as identified by MW-6. Water and free phase product remain exclusively at this low point in the permeable zone to the present 15 years after the massive McKesson dumping was discontinued in 1986. Dehydration of the permeable zone occurred in the years after abatement of the daily discharge of DNAPL solvents and water onto the Angeles property, with residual

DNAPL, LNAPL and water remaining in the sump like deeper configuration identified by MW 6 on the Angeles property.

Such large and long term VOC releases, which have been documented at the McKesson property (See Appendix C), can easily diffuse contaminants upgradient from the source for extended periods of time and result in partitioning in all phases. Since contaminants spread away from a source by diffusion and dissolved VOC concentration remain approximately 5 times higher along the southern side of the property than the northern side, it is concluded that the source of dissolved VOCs is located south of the property. This conclusion will be further clarified as recent groundwater sample results from the McKesson property are identified, possibly with higher dissolved phase concentrations of VOCs.

Furthermore, groundwater test data supports that intrinsic reduction of the DNAPLs flowing from the McKesson site through the permeable shallow soil is most probably occurring effectively due to the stripping of the DNAPL from the soil and water by the LNAPL petroleum constituents, e.g., xylene, toluene. No parent chlorinated compounds such as 1,1,1 TCA, PCE, and TCE were identified in concentrations greater than 100 ppb in the north side monitoring well MW-7 and greater than 130 ppb in the south side monitoring well MW-3. However, the dissolved reduction products 1,1 DCA, 1,1 DCE and cis-1,2 DCE were identified in the Gasper/Hollydale Aquifer on the south side of the former Angeles Chemical Company property at concentrations up to 10,300 ppb and up to 2,670 ppb on the north side (See Table 1).

It is further concluded that co-metabolic reduction is occurring between the LNAPL/DNAPL co-solvents since significant concentrations of 1,1,1 TCA, for example, were identified in the LNAPL collected from MW-6 at up to 2.8% of the free product mixture. However, no significant mass of 1,1,1 TCA was identified in the shallow soil in MW-6 and no evidence of migration of these parent compounds was identified to the Gasper/Hollydale Aquifer in MW-7. It is concluded, therefore, that the stripping by preferential partitioning into the more compatible LNAPL has removed the parent products from the previously saturated shallow soil/groundwater permeable zone from 15' bg to 30' bg. Identification of daughter products, exclusively, in the groundwater at adjacent depth to and in the groundwater down-gradient of MW-6 supports that the parent products are partitioning into the LNAPL which are co-soluble at a much greater rate than water.

8.0) RECOMMENDATIONS

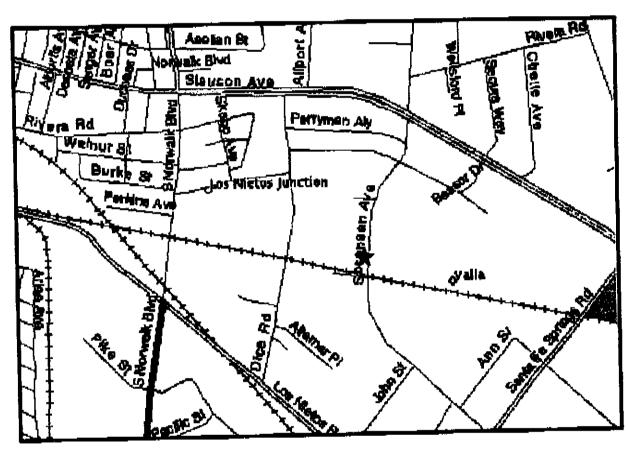
BEII recommends that quarterly groundwater monitoring for VOCs and dissolved

metals be continued at the former Angeles Chemical Co. property.

BEII further recommends that the probable south side source be monitored in dissolved, vapor and free phase near the two property boundaries.

BEII also recommends that a feasibility study of enhanced co-metabolism of the parent VOC products be performed based upon the preliminary data indicating completed first stage co-metabolic reduction of the parent VOCs. Details of the study will be provided in a work plan to be submitted to the DTSC on an agreed date.

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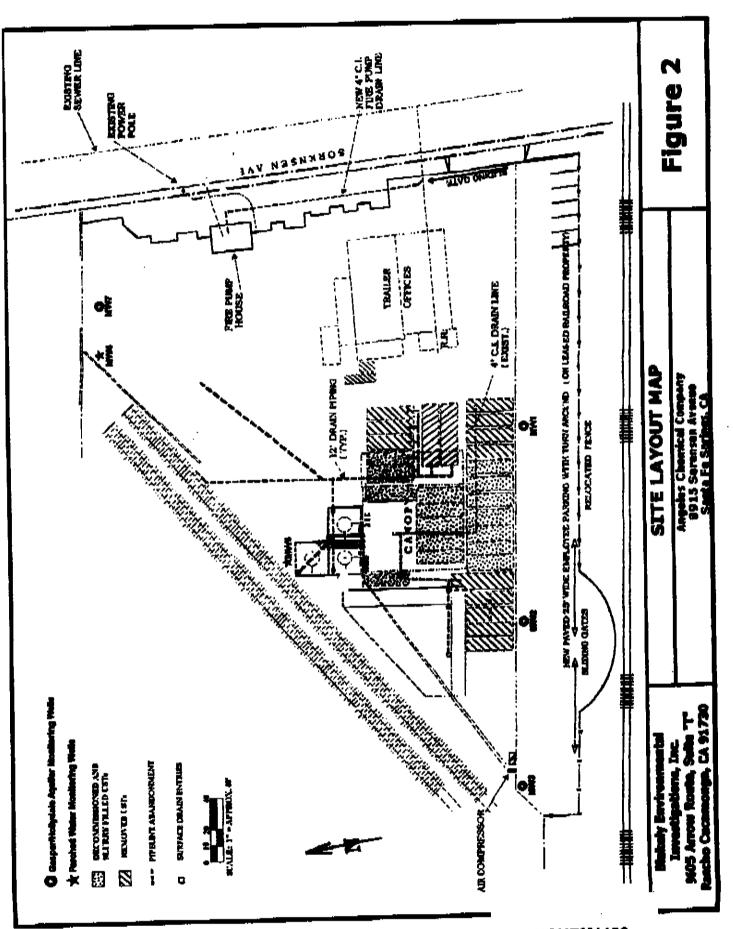
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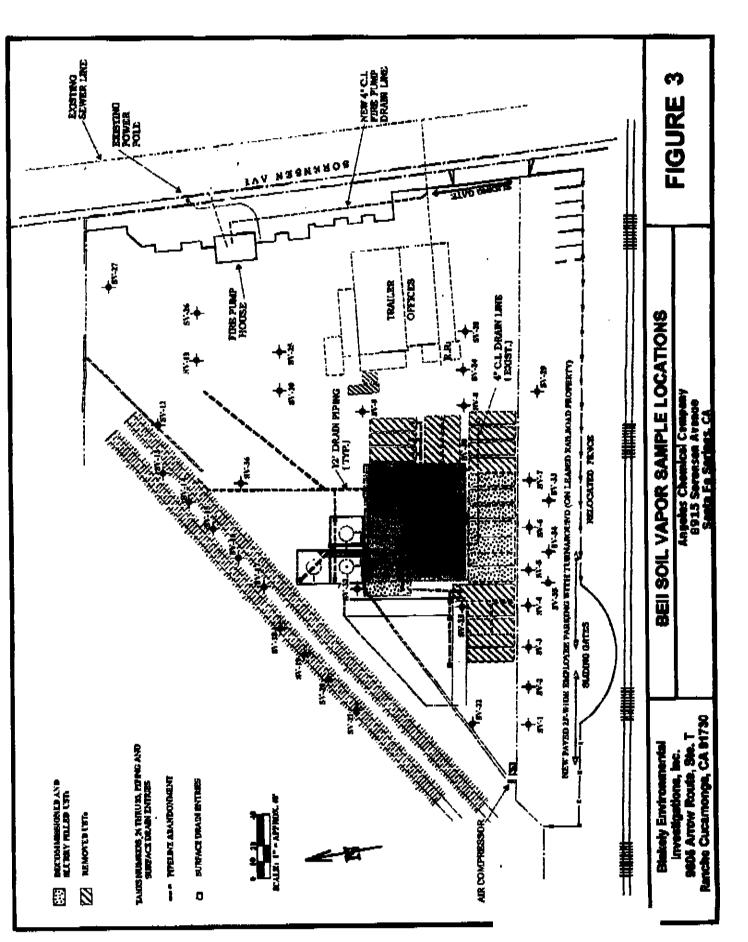


Blakely Environmental Investigations, Inc. 9605 Arrow Route, Suite "T" Rancho Cucamonga, CA 91730

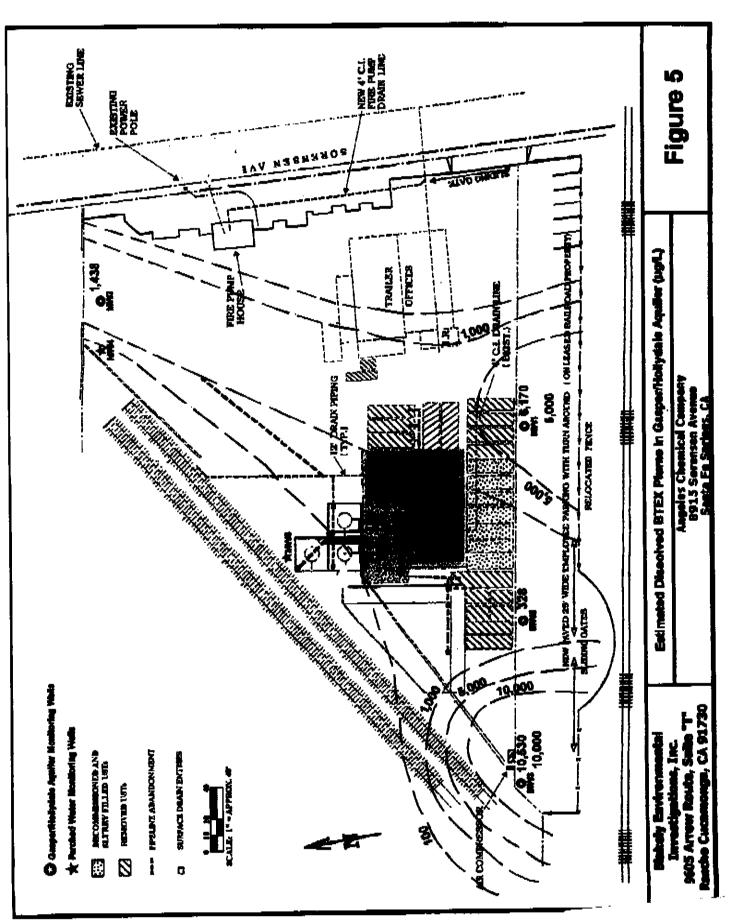
Site Location

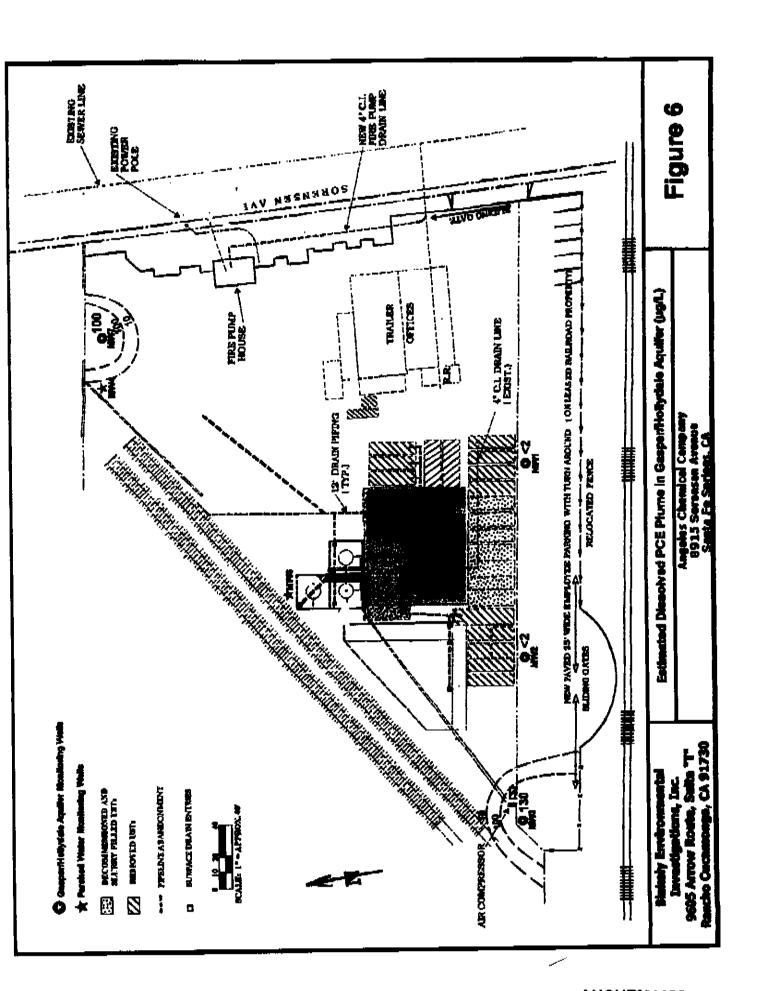
Former Angeles Chemical Comany 8915 Sorensen Avenue Santa Pe Springs, CA FIGURE 1

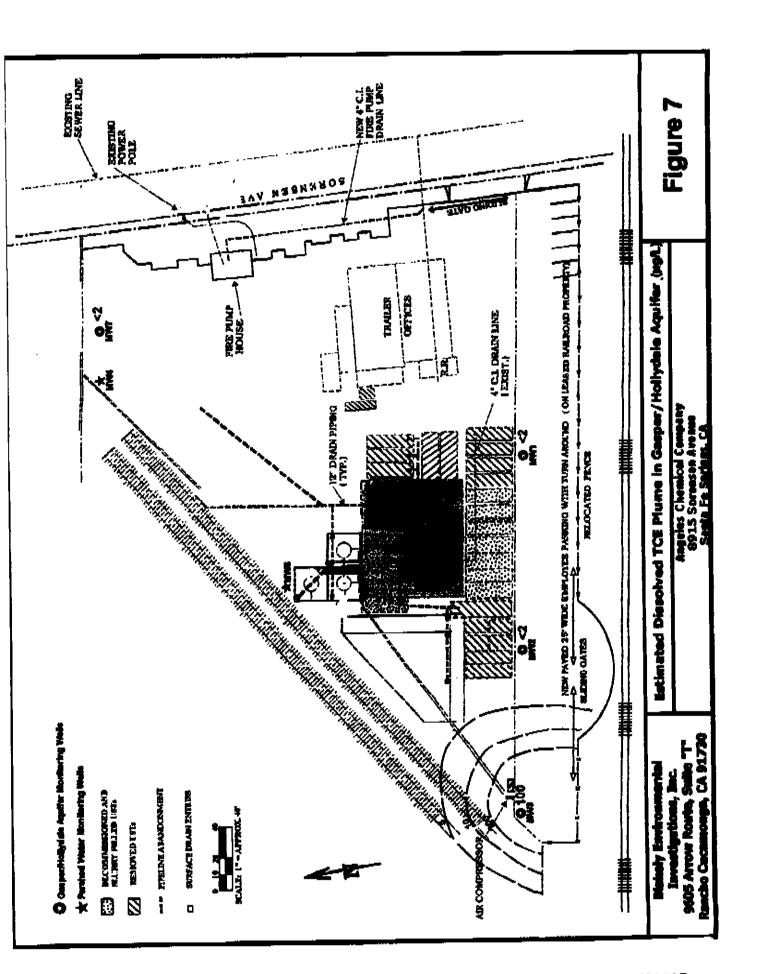


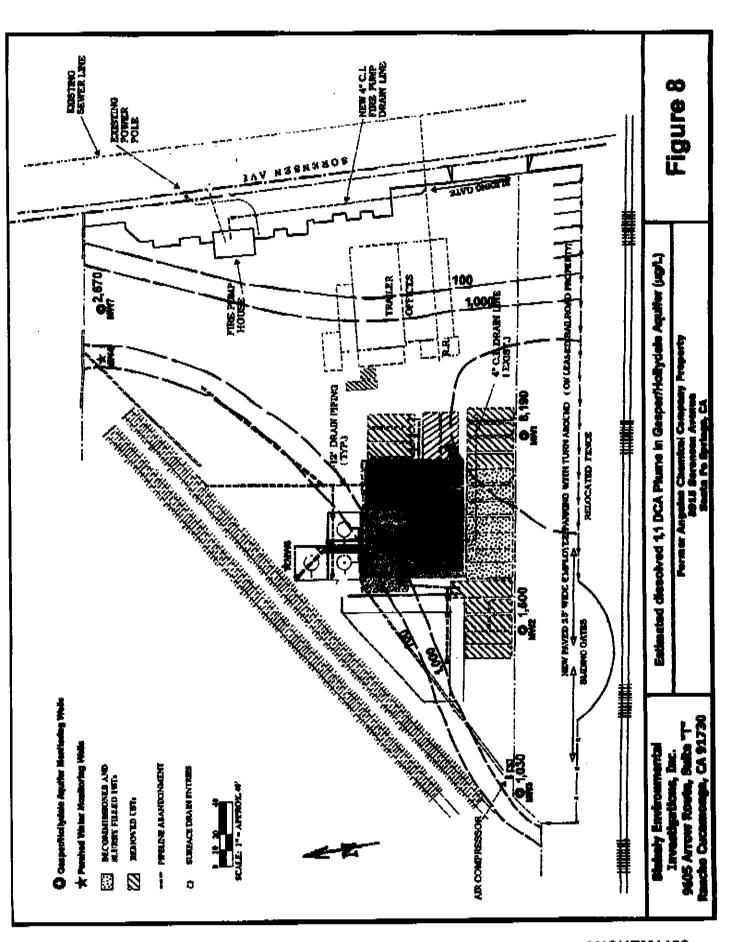


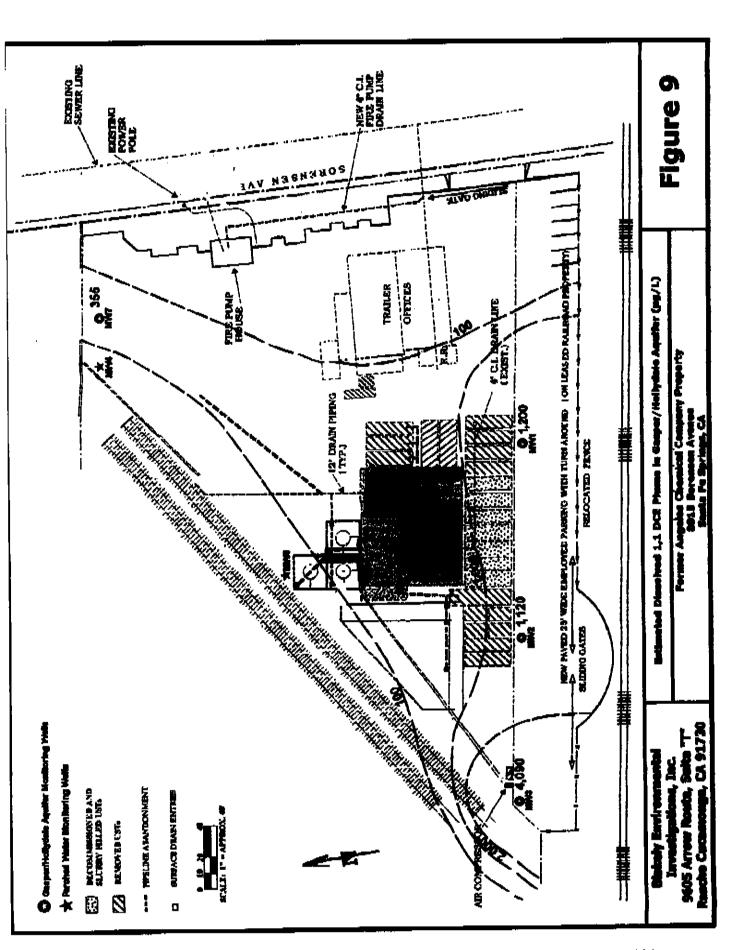
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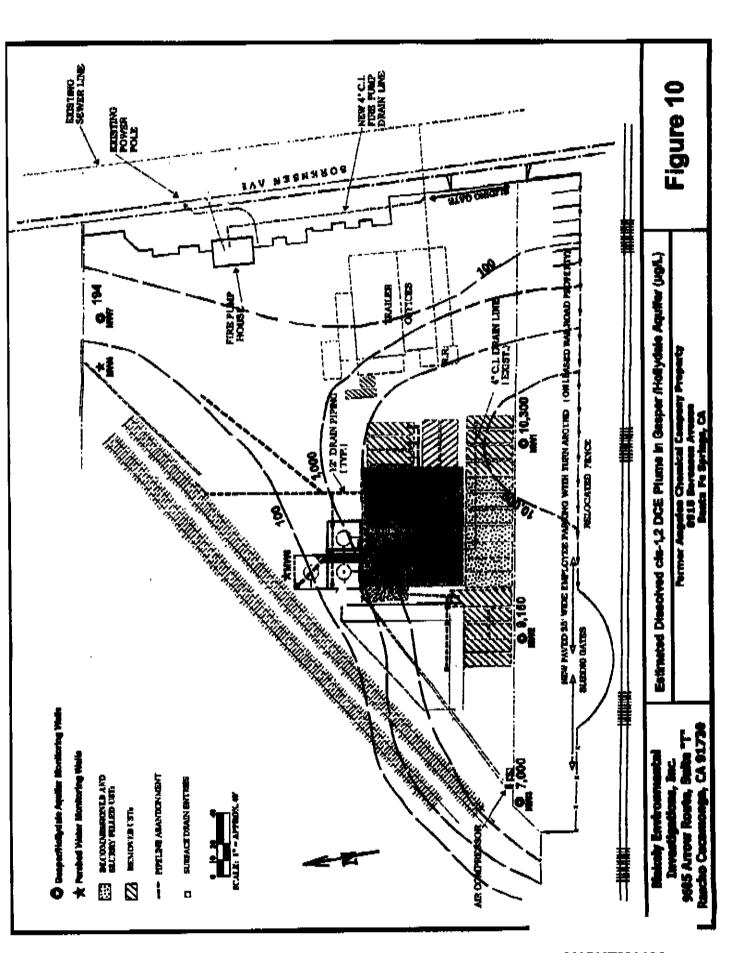


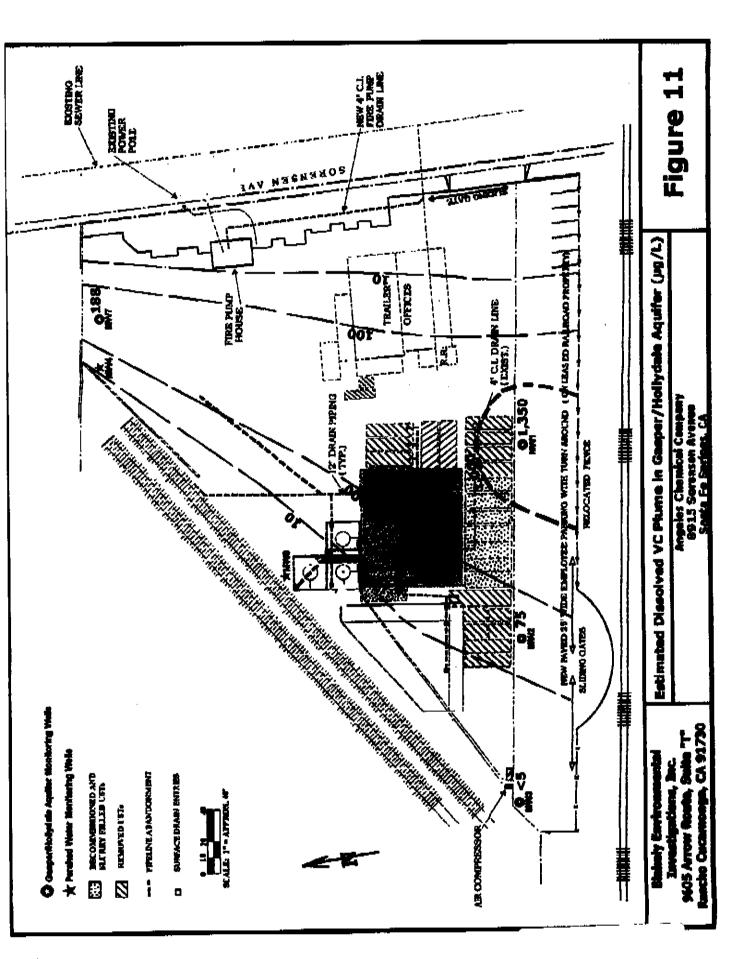


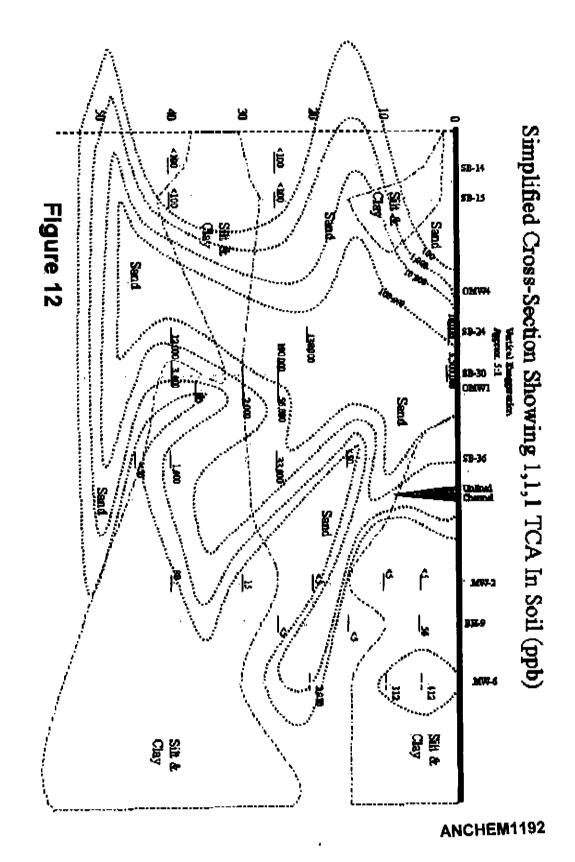












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VOCs	Date	MW-1	MW-2	MW-3	16W-4	MW-6	MW-7
Acetone	Oct-01	<25	<26	<25	NS-NW	<25,000	1,190
		404	<100	65	111	795	46
Benzene	Feb-94	194		73	NS-FP	NS-FP	65
	Nov-00	<2,500	61	110	NS-NW	110,000	55
	Oct-01	125	105	110	140-1444	110,000	- 30
2-Butanone (MEK)	Feb-84	NA.	NA	NA NA	· NA	NA I	NA NA
2-0-04110110 (111211)	Nov-00	3,100	<10,000	<10,000	NS-FP	NS-FP	1,400
	Oot-01	<25	<25	500	NS-NW	<25,000	980
n-Butytbenzene	Oct-01	<5	<5	<5	NB-NW	1,400,000	<5
				 	4-4-4		2 4 20
1,1-Dichloroethane	Feb-94	649	1,130	85	1410	2,260	2,130
	Nov-00	17,000	1,800	800	NS-FP	NS-FP	2,800
	Oat-01	8,190	1,500	1,030	NS-NW	592,000	2,670
1,2-Dichloroethane	Feb-94	<100	<100	<50	<100	1140	31
1,4-DIGINO1060 MING	Nov-00	<2,500	<500	₹500	NS-FP	NS-FP	<500
	Oot-01	<5	3	45	NS-NW	<5,000	<u> </u>
			1 1	 		 -' 	
1,1-Dichloroethene	Feb 94	2,210	2,460	2,800	806	1,240	151
	Nov-00	3,000	<600	2,900	NS-FP	NS-FP	350
	Oct-01	1,200	1,120	4,090	NS-NW	417,000	355
	F-1 04			 	BIA I	NA.	NA.
cis 1,2-Dichloroethene	Feb-94	NA 20.000	NA NA	NA 5 700	NS-FP	NS-FP	210
	Nov-00	20,000	9,500	5,700 7,000	NS-NW	1,000,000	194
	Oct-01	10,300	9,150	1,000	L42-1444	1,000,000	13-7
ans 1.2-Dichloroethene	Feb-94	NA NA	NA	NA	NA I	NA	NA.
ans 1,2-Okasokoconcaic	Nov-00	<2,500	<500	<500	NS-FP	N8-FP	<500
	Oct-01	<6	<5	-55 2	NS-NW	<5,000	<5
				1	110,110	1 3,222	1
Ethylbenzene	Feb-94	333	1,720	115	1,180	1,910	45
	Nov-00	960	120	1,000	N8-FP	NS-FP	82
	Oct-01	806	197	1,550	NS-NW	4,320,000	107
							<u> </u>
isopropylbenzene	Oct-01	<5	<5	<5	NS-NW	636,000	<5
Methylene Chloride	Feb-94	1,220	2,960	6,530	4,760	21,400	<50
Mennyene Chronice	Nov-00	1,100	180	5,600	NS-FP	NS-FP	180
110	Oct-01	<5	45	<5	NS-NW	<5,000	<5
	1				1		
4 Methyl-2-pentanone	Oct-01	<25	<25	4,130	NB-NW	<26,000	625,
Naphthalene	Oct-01	185	76	<5	NS-NW	1,680,000	85
n-Propylberzene	Oot-01	ব	<5	<5	NS-NW	1,870,000	<5
II TI VEYNOREONO			H	+ ~ +	100 100	.,,	1
Tetrachioroethene	Feb-94	662	2,150	5,370	3,320	2,130	134
	Nov-00	<2,500	<500	130	NS-FP	NS-FP	<500
	Oct-01	₹2	<2	130	NS-NW	531,000	100

...

		1	T 1	1	A Method 82		1
VOCs	Date	MW-1	MW-2	MW-3	MW-4	MW-6	MAY-Z
1.1.1-Trichloroethane	Feb-94	9,370	3,470	444	36,200	114,000	90
.,.,.	Nov-00	<2.500	<500	76	NS-FP	NS-FP	<500
	Oct-01	<5	<5	<5	N8-NW	28,100,000	<5
Trichloroethene	Feb-94	7,160	3,040	1,730	14,300	1,320	45
	Nov-00	<2,500	<500	1,500	NS-FP	NS-FP	<500
	Oct-01	~2	<2	100	NS-NW	763,000	~2
2,4-Trimethylbenzene	Oct-01	1,590	18.9	345	NS-NW	22,100,000	200
3,5-Trimethylbenzene	Oct-01	470	62.9	145	NS-NW	5,400,000	25
Toluene	Feb-94	560	7,390	579	12,700	15,300	398
	Nov-00	4,000	57	3,700	NS-FP	NS-FP	600
	Oct-01	2,470	26	5,150	NS-NW	9,010,000	975
Vinyl Chloride	Oat-01	1,350	75	<5	NB-NW	<5,000	188
Xylenes	Feb-94	2,192	7,790	1,014	4,362	4,710	186
	Nov-00	3,400	<500	2,500	NS-FP	NS-FP	247
	Oct-01	2,770	2	3,720	N8-NW	10,370,000	301
DTW	Feb-94	30.05	28.80	29.70	23.36	24.85"	24.53'
	Nov-00	35,62	36.28	36.42	26.20	28.52	28.19
	Oct-01 Nov-01	37.41'	37.91' NA	39.19" NA	26.35° 26.36	NA 28.85'	28.70°
	1404-01	- 1					
Screened Interval		30-50' bg	30'-50' bg	29'-49' bg	17'-27' bg	20'-30' bg	35'-56' bg
TIRL D							
TW= Depth to Water. A= Not Analyzed.	 	 	+	+ +		 	+
S-FP= Not Sempled Fr	e Product	present.					
S-NW= Not Sampled N	<u>lot Enough \</u>	Nater prese	nt.				
ue= Chemicals stored o		1	<u> </u>		+	- 	-
ed= Transformation cor	npounds iro	m chemical	s stored on-al	18 .		+	+

ANCHEM1196

Table 2: Dissolved Metal Sample Results (mg/L)

<u>Dissolved Metals</u> Antimony	EPA Method 7040	Date Oct-01	MW-1 <0.5	MW-2 <0.5	MW-3 <0.5	MW-4 NS-FP	MW-6 NS-FP	<u>₩₩-7</u> <0.5
Arsenic	7060	Oct-01	0.026	0.061	<0.005	NS-FP	NS-FP	0.071
8arium -	7080	Oct-01	<0.5	<0.5	<0.5	N\$-FP	NS-FP	<0.5
Beryllium	7090	Oct-01	<0.05	<0.05	<0.05	NS-FP	NS-FP	<0.05
Cadmium	7130	Oct-01	<0.05	<0.05	<0.05	NS-FP	NS-FP	<0.05
Chromium	7190	Oct-01	<0.1	<0.1	<0.1	NS-FP	NS-FP	<0.1
Cobalt	7200	Oct-01	<0.1	0.12	<0.1	NS-FP	NS-FP	<0.1
Copper	7210	Oct-01	<0.05	<0.05	<0.05	NS-FP	NS-FP	<0.05
Lead	7240	Oct-01	<0.1	<0.1	<0.1	NS-FP	NS-FP	<0.1
Mercury	7471	Oct-01	<0.001	<0.001	<0.001	NS-FP	NS-FP	<0.001
Molybdenum	7480	Oct-01	<0.04	<0.04	<0.04	NS-FP	NS-FP	<0.04
Nickel	7520	Oct-01	<0.1	<0.1	<0.1	NS-FP	NS-FP	<0.1
Selenium	7740	Oct-01	<0.005	<0.005	<0.005	NS-FP	NS-FP	<0.005
Silver	7760	Oct-01	<0.05	<0.05	<0. 0 5	NS-FP	NS-FP	<0.05
Thallium	7840	Oct-01	<0.2	<0.2	<0.2	NS-FP	NS-FP	<0.2
Vanadium	7910	Oct-01	<0.5	<0.5	<0.5	NS-FP	NS-FP	<0.5
Zinc	7950	Od-01	<0.05	<0.05	<0.05	NS-FP	NS-FP	<0.05

NS-FP= Not Sampled Free Product present.

ANCHEM1197

Recycled Stock # Blakley-G-S

WELL GAUGING DATA

Project # <u>& // 03 </u>	P1 Date 10/30/01	Client	Blu 48 4	Ful.
SiteAngeles	Chem			

		T	<u> </u>	Thickness	Volume of				
	Well Size	Sheen /	Depth to	of Immiscible	immiscibles Removed	Depth to water	Depth to well	Survey Point: TOB	
Well ID	(in.)	Odor	Liquid (ft.)	Liquid (ft.)	(ml)	(fl.)	bottom (fl.)	or TOC	
n=3	4/	odor				31.11	25.00	724	
nw-\$	4	Tolure odor				37.41	62.70	47	
ww.2	Ч					37.91	52.z7	24	
w. 4	4				_	26.35	26.65		
AN C	4	5PH	_)	
uw 7	2					2770	51.85	▼ _	
								 	
		<u></u>							
_									
		-					"		
		 		<u> </u>				_	
						 			
			-		<u> </u>				
	<u> </u>	<u>i</u>		<u> </u>	<u>i</u>	1			<u> </u>

laine Tech Services, Inc. 1680 Rogers Ave., San Jose, CA 95112 (408) 573-0555

\. e_L MONIT	ORING DATA Shark 1
Project #: 0//030~ P/	Client: Blackaly Env Augulos elea
	Start Date: (= /30/01
Sampler: 72	Well Diameter: 2 3 6 8
Well I.D.: Man-/	Depth to Water: 37.4/
Total Well Depth: 62.70	A flor:
Before: After:	Before:
Depth to Free Product:	Thickness of Free Product (feet): YSI HACH
Referenced to: evc Grade	D.O. Meter (II 104 5).
Purge Method: Baller Waterra Disposable Baller Peristaltic Extraction Pump Electric Submersible Other	Well Dismeter Multiplier Well Dismes 1995
49.2	014 6- 1.47
Case Volume Specified Volumes Calculated	Volume
Time Temp (°F) pH Cond	Turbidity Gais. Removes
12:50 71.3 6.80 2031	38 G Odor
TO TACK	18 32 "
7 (00 (00)	10 49 4
13:17 71.4 6.89 1879	
	Gallons actually evacuated: 49
Did well dewater? Yes No	
Sampling Time: cher has defer and	Sampling Date. /2/28/6/
Sample I.D.: Mar-/	Laboratory. Stapes
Analyzed for: TPH-G BTEX MTBE TPH	
Equipment Blank I.D.:	Duplicate I.D.:
Analyzed for: TPH-G BTEX MTBE TPH	
D.O. (if req'd): Pre-p	urge: /t Post-pung
ORP (if req'd):	

	L MONFI لــــــــــــــــــــــــــــــــــــ	FORING DATA	A State (
Project #: 0/1030 /	7/	Client: 3/	sely our.	sayers clan
Sampler: PB / JT		Start Date: /	···	
Well I.D.:	<u></u>	Well Diameter	r: 2 3 🗲	0 6 8
Total Well Depth: 62.3	7	Depth to Wate	x: 37.91	
Before: After:	,	Before:		After:
Depth to Free Product:	. <u></u>	Thickness of F	ree Product (fe	et):
Referenced to:	Grade	D.O. Meter (if	req'd):	Y\$1 HACH
Purge Method: Bailer Disposable Bailer <u>Middleberg</u> Electric Submersible	Waterra Peristaltic Extraction Pump Other	Other:	Dispersible Builder Extraction Port Dedicated Tubing	
9.5 (Gals.) X 5 1 Case Volume Specified Vol		Gals.	0.04 4" 0.16 6"	0.65 1.47
Time Temp (°F) pH	Cond.	Turbidity	Gals. Removed	Observations
18:05 7818 7:07	1907	97	10	
Depth to Free Product: Referenced to: Grade D.O. Meter (if req'd): Purge Method: Bailer Bailer Waterra Disposable Bailer Peristaltic Extraction Pump Dedicated Tubing Electric Submersible Other Other: Well Disposes Multiplier 1° 0.04 4° 0.65 2° 0.16 6° 1.47 3° 0.37 Other radius ° 0.163 Time Temp (°F) pH Cond. Turbidity Gals. Removed Observations				
well Dave	tered on	17 gcl		
Did well dewater Ys	No	Gallons actuall	y evacuated:	2
Sampling Time: alout h	as dut on co			
Sample I.D.: mw. 2		Laboratory: 5	eshe sound a	or to cherr
Analyzed for: TPH-G BTEX		Other Clien	t his durin	
Equipment Blank I.D.:	@	Duplicate I.D.:		
Analyzed for: TPH-G BTEX	MTBE TPH-D	Other:		
O.O. (if req'd):	Pre-purge:	mg/L	Post-purge:	™\$/ _L
ORP (if regid):	D	VZ.mr	D	37

		WEI	L MONITO	RING D	<u>ATA SI</u>	HEET_			
				Client:	Nosel	4			
Project #:	01/830			Start Date	: 101	125/0			
Sampler:	<u>P\$</u>			Well Diam				5 8	ĺ
Well I.D.:	<u> 20.5</u>			Depth to		_			
Total Well	Depth:	52.00			YV ZICCI .			fter:	
Before:	A	fter:		Before:				-	
Depth to F	ree Product			Thicknes	s of Fre	e Produc	YS	HACH	1
Referenced	l to:	avc	Grade	D.O. Me					_
Purge Method		W ler P	Vaterra eristaltic ixtraction Pump	Sampling 1	_	Baller Extraction Dedicated	Port		
	Electric Subme	rsible C)ther		Velt Diencis	Multiplier	West Di	oneter Multiplier 0.65	
			- 24-5	Gals.	1" 2" 3"	0.04 0.16 0.37	6" Other	1.47 radius ^{2 *} 0.163	
1 Case Volum		CITIES VOICE	es <u>Calculated \</u>	<u> </u>	idity	Gals. Re	moved	Observations	_
Time	Temp (°F)		Cond.	35	-	8		odor	
11:15	71.7	7,25	1861		<u></u>			U	
1/3/	71.3	7.19	1878	19		16			
7,3,	2300	dew who	40 CZ	7.9º	<u>e</u>	 -			
\———	<u> </u>					 -			
	 					<u> </u>			
	1 1 (ar?)	#CS	No	Gallo	ns actua	lly evacu	nated:	/7	
Did wel	l dewater?	<u>- (</u>		Samp	ling Dat	te: 10	128/0	<u> </u>	
Samplin	g Time: E	Vient has	dutu ou	Labo	ratory:	54 mp 105	spend	ner to chat	
Sample	I.D.: pole	<u>-3</u>	MTBE TPH	u Other	01000	- h1=	Line		_
Analyz	ed for: TP	H-G BIEN	MTBE TPH		icate I.I				
- Equipm	nent Blank	I.D.:	Time						
		H-G BTEX			nıg	/.		ge:	** / _L
D.O. (i	f req'd):		Pre-p			 -	Post-pur		n
ORP (if req'd):		., te-l	ourge:					

_		V	VELL MONIT	ORING DATA	SHEET	
Project #	: 01,0	130-P1		Client: Black	ely Full a	ropolas chem
Sampler:	03/			Start Date:	10/28/01	ropolos chem
Well I.D.				1	: 🙆 3 4	
Total We		51.85		Depth to Water	r: 27.70	
Before:		After:		Before:		After:
Depth to	Free Produ	uct:		Thickness of F	ree Product (fee	et):
Reference		D.	Grade	D.O. Meter (if	req'd):	YSI HACH
Purge Metho	od: Baller Disposable B Middichurg Electric Subn	>	Waterra Peristaltic Extraction Pump Other		Extraction Port Dedicated Tubing	
3. 7.	_(Gais.) X _ neSr		= //./ nes Calculated Vo	Gals. Well Dismon	0.04 4* 0.037 Other	Diamoter Multiplier 0.65 1.47 r radius ^{2 +} 0.163
Time	Temp (°F)	pН	Cond.	Turbidity	Gals. Removed	Observations
11:35	71.5	6.43	2637	٥٩٤٦	3.7 41.40	O DOR Blue
(0:25	72.1	4.77	2732	175	7.4	٧
10:34	71,7	7,00	2633	Q1	11.1	af
(0:37	72.0	7.04	2435	7200	13	
		· · · · · · · · · · · · · · · · · · ·				
Did well	dewater?	Yes (No	Gallons actuall	y evacuated:	13
Sampling	Time: C!	ent has	dup a coc.	Sampling Date	: (0+30-01	_
Sample 1.			· · · · ·			me to chiert
Analyzed	for: TPH	G BTEX	мтве трн-и	Other:	thent his due	in on cor
Equipmen	nt Blank I.	D.:	@ Time	Duplicate I.D.:		
Analyzed	for: TPH-	G BTEX	мтве три-D	Other:		
D.O. (if re	:q'd):		Pre-purge:	m#/L	Post-purge:	™8/ _L
ORP (if re	 eq'd):		¹⁵ re-purge:	mV	Post-purge:	πV

WELL MONITORING DATA SHEET

WELL GAUGING DATA

Project # 011(15-60-1 Date 16/15/01 Client Blakely EW. Jured.

Site 8915 SORENSEN AVE - SANTA Fe Springs

Well ID	Well Size (in.)	Sheen / Odor	Liouid (ft.)	Thickness of Immissible Liquid (fc)	Vulune of Immiscibles Removed (mi)		Depth to well buttom (fL)	Survey Point: TOB or TOC	
MW-4 MW-6	4	Oder	2629	0.07		26.36		TOC	·· ·
May - 6	4	Odar	∂8.3 0	0.55		28.85		18C	
			<u></u>	. <u> </u>		<u> </u>			
	-								
			•						
		ļ <u>.</u>					. <u></u>		·
		· ,	· ·						
							-4"		111
				<u></u>					

Blaine Tech Services, Inc. 1680 Rogers Ave., San Jose, CA 98112 (408) 573-0555



Environmental Laboratories

760: Telegraph Road, Suite L Momento, CA 90640

Phone (323) 666-0726 Fax (323) 668-1509

11-07-2001

Mr. Hiram Garcia Biakely Environmental Investigations, Inc. 9605 Arrow Highway, Suite T Rancho Cucamonga, CA 91730

Project:

Angeles Chemicals

Project Site:

Angeles Chemicals

Sample Date: 10-30-2001 Lab Job No.:

GR110126

Dear Mr. Garcia:

Enclosed please find the analytical report for the sample(s) received by STS Environmental Laboratories on 10-30-2001 and analyzed by the following EPA methods:

EPA 7000 Series for CAM Metals EPA 8260B (VOCs by GC/MS)

All analyses have met the QA/QC criteria of this laboratory.

The sample(s) arrived in good conditions (i.e., chilled, intact) and with a chain of custody record attached.

STS Environmental Laboratory is certified by CA DHS (Certificate Number 1986). Thank you for giving us the opportunity to serve you. Please feel free to call me at (323) 888-0728 if our laboratory can be of further service to you.

Sincerely.

Roger Wang, Ph. D. Laboratory Director

Enclosares

This cover letter is an integral part of this analytical report.



Environmental Laboratories

7801-Telegraph Pond, Suite L Montebello, CA 90640

Phono (323) 859-3728 Fex (323) 898-1509

11-07-2007

Client:

Blakely Environmental Investigations, Inc.

Project:

Lab Job No.:

OR110126

Angeles Chemicals Angeles Chemicals

Date Sampled: Date Received: 10-30-2001 10-30-2001

Project Site: Matrix.

Water

Date Analyzed:

Batch No.:

1031-M1

10-31-2001

CAM Motals (Dissolved) Reporting Units: mg/L (ppm)

			Reporting Unit	MW-2	MW-3	MW-7	Reporting
<u> </u>	EPA	Method		GR:0126-2	GR10126-3	GR10126-4	Limit
Element	Method	Blank	GR10126-1			ND	0.5
ntimony (Sb)	7040	שא	ND	ND	ND		0.005
	70(0	ND	0.026	0.061	ND	0.071	1
Arsenic (As)			ND	ND	D D	ND	0.5
Barium (Ba)	7080	S O		ND	ND	ND	0.05
Beryllium (Be)	7090	ND	ND		ND	ND	0.05
Cadmium (Cd)	7130	ND	ND	/AD		1	0. i
	7190	ND	ND	ND	ND	ND	1
.hromium (Cr)	·	i	ND	0.12	ND	ND	0.1
Cobult (Co)	7200	ND	1	ND	ND	ND	0.05
Copper (Cu)	7210	ND	ND	·		ND	0.1
	7240	ND	ND	ND	ND	<u> </u>	0 001
Lead (Pb)	·	ND	ND	ND	ND	ND	1031
Mercury (Hg)	7471	ND.		- 	1	ND	0.4
Molybdenum	7480	מא	, ND	ND	ND	į ND	·
(Mo)	, , , ,		<u> </u>	ND	ND	ND	0.1
Vickel (Ni)	7520	טא	ND		ND ND	ND	0.005
Scknium (Se)	7740	ND	שא	DM	ł	ND	0.05
·	7760	מע	ND	ND	ND	· ·	· · · · · · · · · · · · · · · · · · ·
Silver (Ag)	·	NU	ND	ND	40	עא	0.2
Thailium (Ti)	7840	·	i	ND	ND	770	0.5
Vanadium (V)	7910	ND	ND		ND	ND	0.05
Zinc (Zn)	7950	ND	ND	ND			

Not Detected (at the specified limit). ND:



Environmental Laboratories

7801 Telegraph Road, Suite L Michieletto CA 90840 Phone (323) 888-0728 Fpz (323) 868-1805

Client: Blakely Environmental Investigations, Inc.

Lab Job No.: GR110126

Date Reported: 11-07-01

Project: Angeles Chemicals

Marrix: Water

Date Sampled: 10-30-01

PA 8268B (VOCs by GC/MS, Page 1 of 2) Reporting Unit: ug/L (ppb)

DATE ANA	- VANTO	16.30	MS, Page 1 -	10-30-01	10-31-01	10-20-01	10-30-01
PREPARATION MI		3030	5030	5030	5030	5030	5030
DILUTION	7770		30	10	25	5	1.
LAB SAMP	4C 1 VA		GK10126-1	GR 10126-2	GR10126-3	GR 10126-4	GR 10126-6
LAB SAMP	TIME BALLY		MW-1	MW-2	MW-3	MW-7	Trip
CLIENT SAME		MB	(+2 +				
COMPOUND	MDL		ND	ND	- 80	ND	ND
ichlorodifiuofometrane	5	ND D	<u> </u>	- COM	- מא	ND	NU
hloromeriane	3		1.350	75	ND T	188	עוא
inyl Chloride	3	עא .	ND -	ND	-טא	- עט	עא
Sromomethane	3		ND T	ND O	+ NO -	ND T	NU
hloroethan#	3	שא	ND -	ND	CIN	80	ND
richlorofluoromethine	5	טא טא	1,200	1,120	4,090	353	ДŊ
, J - Dich loroethene	3		- ND-	ND T	לא -	ND	ND
odomethane	3	ND	עא	- ND	ND	ND	עא
Methylene Chloride	5	ND	שא	1 ND	NO	ND	ND
rans-1,1-Dichloroethere	2	טאַ	8,790	1,500	1,030	2,670	עא
1.1-Dichlorcethane	5	ND	ND	- ND	ND	ND	ND
2,2-Dichloropropane	5	ИD	10.300	9,150	7,000	194	NO
cis-1, Z-Dichloroethene	5	ND		GN	ND	שא	ND.
Sromochloromethane	3	ND	ND	- ND	ND	ND D	H ND
Chloroform	- 5	UN	ND	ND	- עא	 ND -	עא יי
1 Z-Dichloroethans	- 3	מא	ИD	+ ND	ND	שא	ND
1.1.1-1 richloroethane	5	עא	ND	- עא	- ND -	ND ND	- סא
Carbon tetrachionole	3	בא ו	ND	טא טא	ND	ND -	- מא
I. I - Dichloropropent	7-3	נוא	ND	105	110	55.0	H NO
Benzenc	1	ΝD	125	- 103 - D	100	ND.	ND TO
Inchlorosthene	<u> </u>	עא	עא	ND -	NO	ND	ND
T,Z-Dichloropropane	5	טא י		- טא	ND.	- QD	- '412
Bromodich loromethane	3	NU		NO-	ND	ND	ND TO
Dibromomethane	5	טא	· · · · · · · ·	- ND	- ND	ND	ND
trans-1,3-Dichloropropene	3	מא	· 1	- ND	ND	- ND	ND
eis-1.3-Dichloropropene	3	ND	· · · · · · · · · · · · · · · · · · ·	- SID-	NO	- ND	ND
1.1.2-Trichloroethane	5	טא ו		- ND	ND-	- סא	עא ו
COmplete Complete	5	NU		NO-	ND-	- עא	טא -
Dibromochloromethane	5	NE		- UN	ND	- ND -	<u> </u>
2-Chloroethylvinyl ether	5	NU		ND ND	- ND -	- (UN -) -	מא
Bromoform	5	NL		+ ND	- ND	NO	ND
ly propylbenzone	3	N	·	- ND	עא 🕂	90	NU
Bromobenzene	- 3	NI	כוא	1 140			



Environmental Laboratories

7901 Talegraph Road, Suite L Montebello, CA 90540

Phone (323) 866-0728 Fax (383) 848-1909

Client: Blakely Environmental Investigations, Inc.

Project: Angeles Chemicals

Lab Job No.: GR110126

Date Reported: 11-07-0!

Matrix: Water

Date Sampled: 10-30-01

EPA 8260B (VOCs by GC/MS, Page 2 of 2) Reporting Unit: µg/L (ppb)

COMPOUND	MDL	MB	MW-1	MW-I	MW-3.	MW-7	Ιπρ
officere	2	ND	2,470	26	5.150	975	ND
etrachloroethene	1 2	ND	NU	ИD	150	10	ND
2-Dioromoethane(EDB)	1.5	עא	- סא	ND	ND	DM	ND
hiorobenzene	1 5	ND	. אט	NU	ND	ND	ND
.1.1,2-Tetrachloroethan		עא	ND	ND	ND _	שא	ND
hyben2cae	1 2	עא	805	197	1,550	107	עא
(&P-Xylenes		מא	1,900	ND	2,890	227	מא
Xylene	7	מא	870	שא	920	74.0	עא –
AVIENC	- 	- ND	ND	- ND	ND	עא	ND
,1,2,2-1 strachlorosman		- עא	NU	עא	ND	NO	ND
,2,3-1richloropropane	- - - - - - - - - - 	ND -	ND	שא	ND	סא	עא
-Propylbenzene	1 3	ND	ND	ND	ND	טא	ND
-Prepytoenzene	 	. <u>MD</u>	ND	ND	ND D	NU	טא
	- 5	- ND	- 80	מא	ND	ND	ND
-Chlorotoluene	3	- CON	470	62.9	145	25	שא
ع,5-Trimethylbenzene	- 5	ND -	- סא	- מא	ND	מא	<u> ND</u>
ert-Butylbenzene	+	ND ND	1,390	18.9	345	200	שא
,2,4-Trimethylbehzene		+ No	NID	H ND	עא	ND	- סא
iec-Butylbenzene	1	NO NO	ND T	ND	- סא	ND	שא
3-Dichlorobenzene	3	NO	ND T	ND	ND	ND	NO
-18оргоруновиеме		+ ND	ND QN	ND	- <u>ND</u>	ND	עא
,4-Dichlorobenzene	3	i	ND ND	ND ND	ND	NO .	ND
,2-Dichlorobenzene	3	ND	ND -	ND ND	+ ND	ND -	
n-Burylbenzene	,	סא	- ND	CN	- ND	ND	' אט
1,4-Trichiorobenzene	3	NO.	ND.	_			
2-Dibrosne-3-	5	ND	ND	MD	ND	ND	ND
Chloropropane	i		ND	- UND	 ND 	שא	שא
Plexac/ilorobutadiene	5_	ND	183	76	- טא	85	ND
Napr. Malene	5	NU	- ND	עא -	- ND	90	ND
1,2,3-Trichlorobenzene	3	ND	1	NU	1 ND	1,190	NO
Acetone	25	טא	טא יי	ND ND	500	980	- ND
Z-Butanone (MEK)	<u>Z5</u>	עא	טא	שא	4,130	625	70
4-Methyl-2-pentanone	25	עא	ND"		1 3,130 -	ND ND	ND.
Z-Hexancre	25	ND	ND	ND	- ND	- GR	ND
MIBE	7	NE,	ND	טא	1417	1 ×2	×1

MDL-Method Desection Limit; MB-Method Black: NO-Not Detected (below DF × MDL).



Environmental Laboratories

7801 Telegraph Flood, Suite L Montebello, GA 90540 Phone (323) 889-0728 Pax (373) 288-1509

Client: Blakely Environmental Investigations, Inc.

Lab Job No.: GR110126

Date Reported: 11-07-01

Project: Angeles Chemicals

Matrix: Product

Date Sampled: 10-30-01

EPA \$260B (VOCs by GC/MS, Page 1 of 2) Reporting Unit: sng/L (ppm)

DATE ANAL			10-30-01	1		
PREPARATION ME	THOD	5030	5030			
DILUTION FA	CTOR		50			
LAB SAMP			GR10126-5			
CLIENT SAMPI			MW-6			_
COMPOUND	MDL	мв				
	3	ND	ND			
Dichlorodiffuoromethane		ND QN	ND	 		
Chiorometriane	3	ND	ND	├ ──	 	
Vinyl Chlorids		<u>טא</u>	ND O			
Bromomethane	5	ND	HD -	<u> </u>	- 	
hiorcethane	5	<u>- עא</u>	ND			†
Inchlorofkeofomethane	2	ND	417			
I_I-Diéfilorocthene	- 5	ND .	NO NO		- -	
odomethane	1	- סא	ND D			
Methylene Chloride	3		ND ND	 	- 	
rans-1,2-Dichloroethene	5	ND	592		_ 	
1.1-Dichloroethane	3	ND.	ND ND		_	
2,2-Dichloropropane	5	עא	1		_	
cis-1,Z-Dichloroethene	5	עא	1,060	 		
Bromochlorometnane	5	NO	ND	<u> </u>		
Chloroterm	5	עא	ND	<u> </u>		
1.2-Dichloroethane	- 5	- סא	שא			
1.1.1-Trichloroethane	3	עא	28,100			
Carbon tetrachloride	- 5	עא	ND			
1,1-Dichloropropene	3	ND.	ND			
Senzene	2	סא	110	<u> </u>		
Trichlorocthene	- 2	טא	/53			
1.Z-Dichloropropane	- 5	ND	עף			
Bromedichieromethans	3	עא	עא	<u> </u>		<u> </u>
Dibromomethane	- 5	טא	עא			<u> </u>
rans-1,3- Dichloropropene	5	ND.	ַ עאַ			
ets-1,3-Dightoropropens	3	סא	ND			
1.1,2-Trichiorocthane	3	ND	NU			
i,3-Dichioropropene	3	מא	ND			
Dibromochloromethane	3	- עא	ND			
Z-Chloroethy vinyl ether	3	שא	־ טא			
Bromotorm	-3	DN	ND			
Isopropylbenzene	- 3	ND	636	T		
Bromobenzene	 3 -	H ND	ND	1		
2: OHIO CHIZOTO						



Environmental Laboratories

78U1 Telegraph Road, Suite L Monteceso, CA 90540 Phone (323) 588-0738 Fax (323) 588-1509

Client: Blakely Environmental Investigations, Inc.

Project: Angeles Chemicals

Lab Job No.: GRI 10126 Matrix: : Product Date Reported: 11-07-01 Date Sampled: 10-30-01

The excell CVC/Ca by CC/MS. Page 2 of 2) Reporting Unit: mg/L (ppm)

EPA 8200B			Page 7 of 2) Ke	paraing Cities	210/2 (P\$1111)	
COMPOUND	MDL	MB	MW-6			
Loluene	2	עט.	9,010			
Tetrachicrosthene	2	ND .	531			
(Z-Dibromcettiane(EDH)	3	ΝD	שא	1		
Chlorobenzene	5	עא	שא	<u></u>		
1,1,1,Z-Tetrachioroettien	5	2	מא			
Ethylbenzena	2	מא	4,320			
M&P-Xylenes	2	ND	6,950	1		
O-Xylene	2	ND	3,120			
Styrene	5	פל	ND			
1,1,2,2-1 Etrachloroethan	5	ND.	ND		<u> </u>	
1,2,3-Trichloropropane	3	עא	טא			
n-Propyloenzene	5	NO	1,870	<u> </u>		
2-Chiorololuene	5	עא	ND	 		
4-Chicrotoluene	. 5	ND	NU			
1,3,5-Trimcthylbenzene	5	עא	5,400		<u> </u>	
tert-Butylbenzene	55	ND	ND			
1.2,4-1 imethylbonzone	5	ND	22,100		. 	
Sec-Buty/benzene	- 5	עא	ND			
1,3-Dichlorobedzene	3	ND	ND			
p-Ixopropytioluene	- 3	ND	ND		_	<u> </u>
7.4-Dichlorobenzene	3	מא	שא			<u> </u>
1,2-Dichlorobenzene	5	ND	ND	 		
n-Butyloenzene	3	עא	1,400			
1, 2, 4- Trichlorobenzane	5	ND	שא			-
1,2-Dibromo-3-	5	ND	DN	1	1	1
Chloropropane	ì		ND -			
Hexachlorobutadiene	5	ND	1,550	 		
Naphthalene	3	ND	ND ND	 		 -
1,2,3- Trichlorobenzene	5		· · · · · · · · · · · · · · · · · · ·		- 	
Acetone	25	ND	עא			
2-Butanone (MEK)	25	טא	ND			-}
4-Methyl-2-pentatione	25	עא	עא			
Z-Hexanone	25	ND.	ND.			
MIBE	- 2	ND	ND			

ND5, "Method Detection Limit; MS=Method Blank: ND=Not Detected (below DF × MDL).



Environmental Laboratories

7801 Talegraph Road, Suite L. Montebello, CA 90640

Phone (323) 866-0726 Fex (323) 866-1509

11-07-2001

EPA 8260B Batch QA/QC Report

Client:

Blakely Environmental Investigations, Inc.

Lab Job No.:

GR110126

Project:

Angeles Chemicals

ST1030-1

Matrix:

Soil

Sample ID: Date Analyzed:

10-30-2001

Batch No:

1030-VOCA

I. MS/MSD Report

Unit: ppb

Compound	Sample Conc.	Spike Conc.	MS	MSD	MS %Rec.	MSD %Rec.	% RPD	%RPD Accept. Limit	%Rec Accept. Limit
1,1- Dichloroethene	ND	20	19.3	17.4	96.5	87.0	10.4	30	70-130
Benzene	ND	20	19.5	19.9	97.5	99.5	2.0	30	70-130
Trichloro-	ND	20	20.6	20.4	103.0	102.0	1.0	30	70-130
Toluens	ND	20	20.9	20.8	104.5	104.0	0.5	30	70-130
Chlorobenzene		20	21 3	20.6	106.5	103.0	3.3	30	70-130

IL LCS Result Unitz ppb

Compound	LCS Report Value	True Value	Rec.%	Accept. Limit
1,1-Dichloroethene		50	87.8	80-120
Benzene	49.2	50	98.4	80-120
Trichloro-ethene	51.1	50	102.2	80-120
	51.4	50	102.8	80-120
Toluene		50	103.0	80-120
Chlorobensens	51.5	318	105.0	

ND: Not Detected (at the specified limit)

14				TOTAL THOUSE							3	TAN AND INSTRUMENT	
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<u>.</u>			1509		istribution	WHIT	E with A	rpar, P	TMK to	courier.			
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ecology and environment, inc.

160 SPEAR STREET, SAN FRANCISCO, CALIFORNIA 94105, TEL. 415/777-2811

Americal Specialists in the Embana.

SCREENING SITE INSPECTION REASSESSMENT

SUMMITTED TO:

Paul LaCourreye, Site Assessment Henager

EPA Region IX

PREPARED BY:

Helena Brykers, Ecology and Environment, Inc. 475

TEROUGE :

Karen Ladd, Ecology and Environment, Inc.

DATE

June 15, 1990

SITE:

Foremost-McKesson, Inc. McKesson Chemical Company 9005 Sorensen Avenue

Santa Pe Springs, CA 90670

TDD#:

P9-9002-014

EPA ID#:

CAD060395753

PROGRAM ACCOUNTS: PCA0243SAA

FIT REVIEW/CONCURRENCE:

Keron Lad 6/18/90

cc: FIT Master File

Don Plain, California Department of Health Services

INTRODUCTION

The U.S. Environmental Protection Agency, Region IX, has tasked Ecology and Environment, Inc.'s Field Investigation Team (FIT) to reassess all sites with completed Screening Site Inspections (SSI) in the Comprehensive Environmental Response, Compensation, and Liability Information System (CERCLIS) database that are still being considered for further action. The strategy for determining whether these SSIs actually merit further action is based primarily on each site's potential to achieve a score high enough on the proposed revised Hazard Ranking System (rHRS) for inclusion on the National Priorities List (NPL). This strategy is intended to identify those sites posing the highest relative risk to human health or the environment. All other sites needing remedial or enforcement follow-up will be referred to the states or an appropriate federal authority. Actions and involvement by authorities other than the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) vill also be considered.

hb/fm/si-re

SUPPLANT

The Foremost-McKesson, Inc., site is located at 9005 Sorensen Avenue in Santa Fe Springs, Los Angeles County, California. It occupies 3 acres of land in an industrialized area, south of the Southern Pacific railroad tracks (1).

Foremost-McKesson operated a chemical reformulation and distribution facility at the site from 1976 to 1986. Virgin chemicals were stored in 20 underground tanks and an aboveground tank farm. Two vaste streams were generated at the facility. Corrosive wastes from drum rinsing operations were pumped, at a rate of 1,500 gallons per day, to an on-site neutralisation pit, and later discharged to the los Angeles County sanitary sever. The 55-gallon drums were then sent to a drum recycler. Isopropyl alcohol was used to flush the solvent lines. The vastes were stored in drums in a designated hazardous waste storage area on site and later disposed of off site (1).

Foremost-HcKesson has not operated at the site since 1986. The California Department of Health Services (DOHS) is currently overseeing the closure of the facility. The tanks, underground and aboveground have been emptied, and the contents were disposed of off site. The drums have been transported off site (2).

Several apparent problems can be identified at the site. A 10,000-gallon spill of sulfuric acid occurred in 1979 in the tank farm area as the result of a faulty tank valve. Additional chemical spills occurred in the tank farm area in 1977, 1978, and 1980. Hovever, the specific chemicals involved are not known to FIT. Although the tank farm area is bermed, it sits on gravel rather than pavement. Results of 1984 DOHS soil sampling efforts indicated that surface soils in the tank farm area vere contaminated with up to 33 parts per million (ppm) tetrachloroethene. In 1985, DOHS soil sampling in the same area detected 100 milligrams per kilogram (mg/kg) of acetone. Also, trichloroethane, methyl ethyl ketone, p-dioxane, tetrachloroethane, and xylene were detected in soil samples, however the concentrations are unknown to FIT (1). The results of a 1989 DOHS sampling effort indicate that trichloroethane at 0.11 mg/kg and tetrachloroethene at 0.5 mg/kg are still present in surface soils (2,4). In 1980, the Los Angeles County Engineers issued a violation to Foremost-McKesson for an observed discharge of industrial waste to an unlined ditch behind the property. The ditch receives surface water runoff from the site via a 2-foot by 2-foot by 4-foot concrete catch basin (1).

The site is located in the Montebello Forebay area of the Central Basin, which is located in the Coastal Plain of Los Angeles County. The area is underlain in order of increasing depth by the Artesia, Gardena, Lynwood, Silverado and Sunnyside aquifers. According to a California Department of Vater Resources report, these aquifers consists of mostly clay and gravel and are interconnected within 2 miles of the site (5).

The Bellflower aquitard, which exists just below ground surface, is approximately 20 to 40 feet thick and pinches out within 2 miles northwest of the site. It consists of clay and sandy clay. However,

percolation throughout the Bellflover aquitard is high, rendering it ineffective as a barrier to flow (5).

The nearest drinking vater vell is located 0.33 miles north of the site (1). It is perforated at 200 to 288 feet and 300 to 900 feet in the Silverado and Sunnyside aquifers (6). The vell is part of an interconnected system of vells in the Santa Fe main system which serves a predominantly commercial area. During the day, approximatly 100,000 workers and residents use the drinking vater, and at night approximately 10,000 residents use the vater (7). A City of Norvalk vell is approximately 2.5 miles south of the site, which serves 2,200 people (8). Southern California Water Company has a vell approximately 2 miles south of the site, which serves a population of 7,000 (9). Both of these vells tap the Silverado and Sunnyside aquifers. The total population served by vells within 4 miles of the site is approximately 109,200.

Groundwater flow is toward the southeast (1). The annual net precipitation is 3.15 inches (10,11).

In 1989, TCE was detected at 2.8 milligrams per liter (mg/l), in the nearest Santa Fe Springs municipal well, 0.33 miles hydraulically upgradient of the site, by the Sante Fe Springs Department of Public Works. Well testing three months later detected no contamination present (12).

The potential for release to groundwater is high, given inadequate containment of hazardous substances, a depth to groundwater of 57 feet bgs, and the moderate permeability of geologic materials in the area-Some of the hazardous substances on site that are, or were, available for release to groundwater, and are highly toxic and mobile in groundwater, include methyl ethyl ketone, p-dioxane, acetone, trichloroethane, and toluene.

Storm runoff from the site flows to a concrete catch basin (2-foot by 2-foot by 4-foot), which has a locking gate valve to control release to an unlined ditch which parallels the northern border of the property. This unnamed ditch drains into the Sorensen Avenue Drain approximately 0.5 miles from the site, and in 2.5 miles it joins La Canada Verde Creek. After approximately 5 miles downstream, it merges with Coyote Creek. In another 6 miles downstream, Coyote Creek empties into the San Gabriel River (13,14). The 2-year, 24-hour rainfall in this area is 3 inches (15).

A potential exists for a release to surface vater based on inadequate containment and the proximity of the site to surface vater. Hazardous substances available for a release to surface vater include acetone and trichloroethane, which are highly toxic and are persistant in rivers (2).

The facility claimed that it applied for an National Pollutant Discharge Elimination System (NPDES) permit, however according to the NPDES database, it never received a permit (1,16).

Although a potential for a release to surface water exists, there are no drinking water uses along the surface water pathway within 15 miles

hb/fm/si-re

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DEPARTMENT OF MEATH SERVICES
TO SOUTH MONUMAY, ROOM 1128
TO AMERICA CA 40017

(213) 620-2380

October 5, 1984

Douglas L. Eisner HeKesson Chemical Company One Post Street San Francisco, CA 94104

Dear Mr. Eisner:

EFA+ CAD 060395753

On October 3, 1984, Megan Robinson, DOMS representative, met with Dudght Landry and yourself to discuss cleanup plans of the contaminated soil at the Santa Pe Springs Facility. Although there is no definitive cleanup standard for unlawfully disposed hazardous waste, the Department's general standard is that all such waste must be removed and lawfully redisposed. As an interim measure, you are to remove surface and near-surface contamination to the levels detailed in the California Assessment Manual (CAM) for those components for which CAM standards are listed.

To determine the full nature and extent of contamination, you are to submit a site characterization and cleanup plan to this office by October 26, 1984. Such plan will outline McKesson's proposal with regard to complete identification of surface and subsurface contamination. The plan must propose methods of identification of groundwater contamination associated with the unlawful disposal. Additionally, include any background information regarding all spill incidents and material stored in the now removed storage tank.

For assistance with the site characterization and clean-up, contact Mestor Acedera at this office.

Your compliance with the requirement for a site characterization and cleanup plan in no way releases you from applicable penalities.

If you have any questions, please do not hesitate to call.

Sincerely.

Jid Smith, Manager

Complaints & Investigation

Southern California Section

Toxic Substances Control Division

JS : #W

cc: Dwight Landry McKesson Chemical Company

ALIFORNIA REGIONAL WATER QUALITY CONTROL BOARD-

OS ANGELES REGION

M SOUTH THO LOWAY, SUITE HIST SE COSTA, CAUSERNA CANT

J) 520-4400

March 7 1981

NoRes of Chemical Company

8005 i tursen Avenue

Sant: 1: Springs, CA 90638

Alba: Mr. Don Wettstead, Bulk Plant Manager

No: Waste Discharge (B VI a-62)

Centlemen:

Nour August 22, 1980, letter to this Board stated that all vustewater accumulated in dikad areas at your plant is now neutralized and sewered, and that you therefore found it unnecessary to maintain your NPDES permit. On that basis we allowed your NFDES permit to expire on November 30, 1980.

On January 22, 1981, a routine inspection of your facility by this Board's staff, together with Mr. Don Wettstead, discharged that washdown wastewaters from the container preparation and filling sheds were being discharged to a drainage ditch.

This discharge is not in accordance with the California Water Coie.

Mr. Wennstead equand to pump the westewaters from the ditch to the similary sever, to remove soils for disposal at a Class I disposal site, and to take steps to prevent the discharge of wastewater from the property in the future.

You are requested to provide a written report to this Board by April 7, 1931 of the corrective measures your company will take to preclude such waste discharges in the future and the timetable for implementation of those measures.



COUNTY OF LOS ANGELES DEPARTMENT OF COUNTY ENGINEER PROJECT PLANNING AND POLLUTION CONTROL DIVISION

INSPECTOR'S REPORT

TO: CANL SLOBERG DATE: Avery 28, 1980
FROM: 1006 FILE: 1-2130-14
INDUSTRY: ME KESSON CHEMICAL CO.
9005 S. SORENSON AVE, SOUTH TE SPENGS
REPORT: DN A VIOLATION CORRECTION ON July 15, 1980
THE FOLLOWING VIOLATION WERE CORRECTED:
1. REPAIRED THE KEY SWITCH PUMP ATTHE
INDUSTRIAL WASTE TREATMENT SUMP.
2. Clemical Spile was clean-up at tite
TANK FARM ALEA.
3. INDUSTRIAL WASTE DISCHARGE RECORDS AND
KEPT ON FILE FOR INSPECTION.
THE FOLLOWING VIOLATION REMAINS:
L REMOVED CONTAMINATED SOIL TO A
LEGAL FOINT OF DISPOSAL
2. KEEPING YAND CATCH BASIN SUMP FRE
OF CHOUCAL WASTE.
A FOLLOW UP INSPECTION ON AUGUST 26,1980,
I FOUND THE CONTAMINATED SOIL HAS TREEN
REMOVED TO THE TSKK DUMP BUT THE CHAMICAL
REMOVED TO THE TOKK DUMP BUT THE CHANICAL WASTERN THE YAND CATCH BASIN, AND THE CHANICAL
IS DESTRUCT OUT TO THE UNLINE DITCH A NOTICE
F VIOLATION AND ARDER TO COMPLY ISSUED TO
MR. DOW WETTSTEAD, PLANT MANAGER.

CITY OF STATE SPRULS

NOTICE OF VIOLATION AND ORDER TO COMPLY

Date Aulust 27,1980

TO ME KESSON CHONICAL M.-ATTENTION: MR. DON WETTSTEAD LOCATION 9005 5. SORENSEN AVE., SANTA FE SPRINGS

You are hereby directed to correct the following violations of City of Santa To Spends Ordinance No. Stor. and/or the conditions and limitations of Industrial Waste Disposal Permit No. Stored by Section 1980.

DIECHARGIALA INDUSTRIAL WASTE INTO THE UNIVERDICH IS AN VIOLATION OF THE ABOVE CITY ORDINANCE.

YOU ARE HOREBY DIRECTED TO CASE AND DESIST
THE DECHARGE, AND CLEW ID THE UNLINE DICH AT ONCE.

YOU ARE ALSO DRECTED TO SUBMIT A WRITTEN.

REPORT TO THE OFFICE BY SEPTEMBER 10, 1980.

PROVIDING A COMPLETE DETAILS OF THE ACTION TOKEN.

TO COMPLY WITH THE ABOVE VIOLATION.

SHOULD YOU HAVE ANY RUFFRION, PLEASE CAN
HE AT SUG-7011, EXTENDION 255 BETWEEN THE

DEPT OF COUNTY ENGINEER

SWITATION DIVISION

14623 BELFLOWER BLID.

BELFLOWER, CA. 90706

COUNTY OF LOS ANGELES DEPARTMENT OF COUNTY ENGINEER PROJECT PLANNING AND POLLUTION CONTROL DIVISION

INSPECTOR'S REPORT

TO: DATE: DATE:
FROM: 1 FILE: I-2/30-1#
INDUSTRY: MEKESSON CHEMICAL COMPANY
- gas S. Sozassen ME. Sonta To Spring
REPORT: ON A POSTINE INSPECTION ON 3/4/80, I
NOTES THE FOLLOWING VICLATION.
1: A Backen Kar Switch Pump For
purpose of western to the Senter.
2. astonical Spills IN THE Tonk Famous
And alse NOT Closed Up.
3. No INDUSTRIAL WASTE DISCHARGE RELOAD,
WHICH REDUINED BY PANIT CONDITIONS.
4. ACID OF PHI FOUND IN THE YAMO.
CATCH BASINI
ME BRION BEAMEN, ASSISTANT MANUSCE OF
THE ABOUT COMPANY WAS INSTRUCTED TO Comply
By Apa. 6 7, 1980.
RE-INSPECTION ON April 7, 1980 ALL THE
ARME VIOLATION WAS NOT CONFERENCE, AND A
NOTICE OF VIOLATION AND MOON TO Comply 15500
To THE Congany, (Notice ATTACHOS)
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EDMUND G. BROWN JR., Gararte

STA OF CASHORNIA - RESOURCES AGENCY

CALIFORNIA REGIONAL WATER QL..LITY CONTROL BOARD. LOS ANGELES REGION

TOUCH BROADWAY, SUITE 4027 3 ANGELES, CALIFORNIA 70012 (213) 620-4460

McKesson Chemical Company

9005 Sorensen Avenue

Santa Fe Springs, California 90670

Attention: Mr. E. W. Boyd, Bulk Plant Manager

Re: Waste Discharge Requirements (CA0057631)(CI 6213)

Gentlemen:

Reference is made to Order No. 75-151 (NPDES Permit No. CACO57631), prescribing requirements for the disposal of wastes from your facility at the above location.

Effluent Limitation A-7 provides that wastes discharged shall not contain visible oil or grease, and shall not cause the appearance of grease, oil or oily slick, or foam in the receiving waters or on channel banks, walls, inverts or other structures.

On December 27, 1977, this Board's staff found that your discharge caused considerable foaming in the unlined ditch tributary to North Fork Coyote Creek, in non-compliance with the above-cited requirement. The fosming was traced to have started from open plant areas near your filling and container preparation sheds.

The problem was discussed with your Mr. Brian Beamer, who was requested to take immediate corrective and preventive measures to assure that the discharge of wastes fully comply with requirements. It was pointed out to Mr. Beamer that you might consider such measures as constructing rolling berms around the filling and container preparation sheds, and/or installing a water-tight valve in your yard sump, instead of the drop gate valve you have currently in use. The latter has already caused you problems when we found an illegal discharge from it on December 2, 1977, as covered by our letter dated December 13, 1977.

It is requested that you submit a written report to this Board by January 20, 1978, providing complete details of the corrective and preventive measures you will take (or have already taken), to bring the discharge of wastes in full compliance with requirements, and the timetable thereof.

If you have any questions concerning the above, please contact Mr. Miller E. Chambers at the above number.

Very truly yours,

RAYMOND M. HERTEL Executive Officer

cc: State Water Resources Control Board, Legal Division

Attn: Mr. Harry M. Schueller
Los Angeles County Engineer, Project Planning and Pollution

Control Division
Los Angeles County Flood Control District

Park - It you with

STATE OF CALIFORNIA-RESOURCES AGENCY

CALIFORNIA REGIONAL WATER QUALITY CONTROL BOARD— LOS ANGELES REGION

SQUIH BROADWAY, SUITE 4027
... ANGELES, CAUPORNIA 90012
620-4460

DEC 131977

McKesson Chemical Company 9005 Sorensen Avenue Santa Fe Springs, California 90670

ATTENTION: Mr. E. W. Boyd, Bulk Plant Marager

RE: Waste Discharge Requirements (CACO57631) (CI 6213)

Gentlemen:

Reference is made to Order No. 75-151 (NPDES Permit No. CAOO57631), prescribing requirements for the disposal of wastes from your facility at the above location.

Effluent Limitation A-7 provides that wastes discharged shall be limited to rainwater runoff during and immediately after rainfall only, as proposed.

On December 2, 1977, this Board's staff found that you had a discharge from your yard sump. Apparently, the discharge resulted from the sump valve not being tightly closed. This discharge is in violation of the above-referenced requirement.

In addition, we also observed that the sump contained a substantial amount of oil.

The problem was discussed with your Mr. E. W. Boyd, who was requested to have the sump cleaned up and the valve tightly closed so that there is no discharge except as authorized by and in compliance with the waste discharge requirements. Mr. Boyd promised to comply with the request.

It is requested that you submit a written report to this Board by December 28,1977, providing details of the corrective measures you will take (or have already taken), including additional steps necessary to assure that similar incidents are not repeated. Please note that the yard sump must be kept clean at all times so that in event of rain, there shall be no discharge of oil or other pollutants except in compliance with discharge requirements.

If you have any questions concerning the above, please contact Mr. Miller E. Chambers of this Board's staff.

Very truly yours.

RAYMOND M. HERTEL Executive Officer

cc: State Water Resources Control Board, Lagal Division

M. Hertel

Attention: Harry Schueller
Los Angeles County Engineer, Project Flanning & Follution Control Division